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TERMINAL (ENTER 1, 2, 3, OR ?):2

* * * * * * * * * * Welcome to STN International * * * * * * * * * *

NEWS	1		Web Page for STN Seminar Schedule - N. America
NEWS	2	JAN 02	STN pricing information for 2008 now available
NEWS	3	JAN 16	CAS patent coverage enhanced to include exemplified prophetic substances
NEWS	4	JAN 28	USPATFULL, USPAT2, and USPATOLD enhanced with new custom IPC display formats
NEWS	5	JAN 28	MARPAT searching enhanced
NEWS	6	JAN 28	USGENE now provides USPTO sequence data within 3 days of publication
NEWS	7	JAN 28	TOXCENTER enhanced with reloaded MEDLINE segment
NEWS	8	JAN 28	MEDLINE and LMEDLINE reloaded with enhancements
NEWS	9	FEB 08	STN Express, Version 8.3, now available
NEWS	10	FEB 20	PCI now available as a replacement to DPCI
NEWS	11	FEB 25	IFIREF reloaded with enhancements
NEWS	12	FEB 25	IMSPRODUCT reloaded with enhancements
NEWS	13	FEB 29	WPINDEX/WPIDS/WPIX enhanced with ECLA and current U.S. National Patent Classification
NEWS	14	MAR 31	IFICDB, IFIPAT, and IFIUDB enhanced with new custom IPC display formats
NEWS	15	MAR 31	CAS REGISTRY enhanced with additional experimental spectra
NEWS	16	MAR 31	CA/CAplus and CASREACT patent number format for U.S. applications updated
NEWS	17	MAR 31	LPCI now available as a replacement to LDPCI
NEWS	18	MAR 31	EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NEWS	19	APR 04	STN AnaVist, Version 1, to be discontinued
NEWS	20	APR 15	WPIDS, WPINDEX, and WPIX enhanced with new predefined hit display formats

NEWS EXPRESS FEBRUARY 08 CURRENT WINDOWS VERSION IS V8.3,
AND CURRENT DISCOVER FILE IS DATED 20 FEBRUARY 2008

NEWS HOURS	STN Operating Hours Plus Help Desk Availability
NEWS LOGIN	Welcome Banner and News Items
NEWS IPC8	For general information regarding STN implementation of IPC 8

Enter NEWS followed by the item number or name to see news on that specific topic.

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result in loss of user privileges and other penalties.

* * * * * * * * * * * * * * * * STN Columbus * * * * * * * * * * * * * * * *

FILE 'HOME' ENTERED AT 13:15:38 ON 15 APR 2008

=> file reg
COST IN U.S. DOLLARS

	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'REGISTRY' ENTERED AT 13:15:45 ON 15 APR 2008
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Property values tagged with IC are from the ZIC/VINITI data file
provided by InfoChem.

STRUCTURE FILE UPDATES: 14 APR 2008 HIGHEST RN 1014671-54-5
DICTIONARY FILE UPDATES: 14 APR 2008 HIGHEST RN 1014671-54-5

New CAS Information Use Policies, enter HELP USAGETERMS for details.

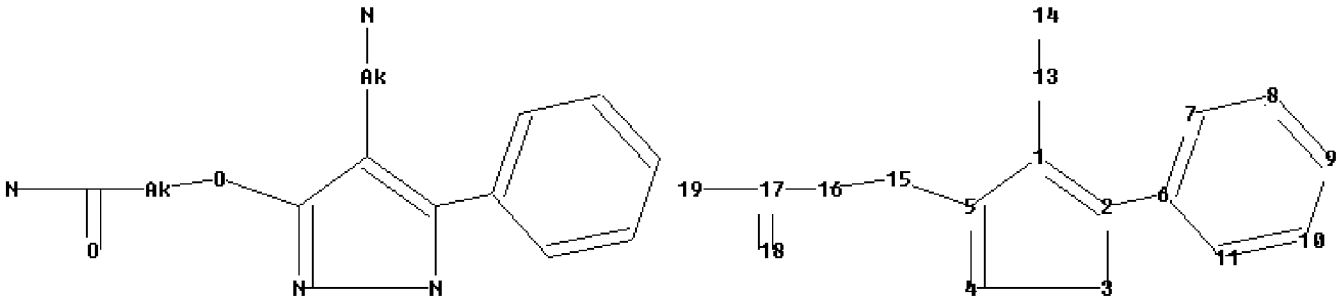
TSCA INFORMATION NOW CURRENT THROUGH January 9, 2008.

Please note that search-term pricing does apply when
conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and
predicted properties as well as tags indicating availability of
experimental property data in the original document. For information
on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=>
Uploading C:\Program Files\Stnexp\Queries\10524977_2.str



chain nodes :
13 14 15 16 17 18
ring nodes :
1 2 3 4 5 6 7 8 9 10 11 19
chain bonds :
1-13 2-6 5-15 13-14 15-16 16-17 17-18 17-19
ring bonds :
1-2 1-5 2-3 3-4 4-5 6-7 6-11 7-8 8-9 9-10 10-11
exact/norm bonds :
1-13 2-3 3-4 4-5 5-15 13-14 15-16 16-17 17-18 17-19

```
exact bonds :
1-2  1-5  2-6
normalized bonds :
6-7  6-11  7-8  8-9  9-10  10-11
isolated ring systems :
containing 1 : 6 :


Match level :
1:Atom  2:Atom  3:Atom  4:Atom  5:Atom  6:Atom  7:Atom  8:Atom  9:Atom  10:Atom
11:Atom  13:CLASS  14:CLASS  15:CLASS  16:CLASS  17:CLASS  18:CLASS  19:Atom



L1      STRUCTURE UPLOADED

=>  d L!
L1 HAS NO ANSWERS
'L! ' IS NOT A VALID STRUCTURE FORMAT KEYWORD
Structure Formats
SIA ----- Structure Image, Attributes, and map table if it contains
          data.  (Default)
SIM ----- Structure IMage.
SAT ----- Structure ATtributes and map table if it contains data.
SCT ----- Structure Connection Table and map table if it contains
          data.
SDA ----- All Structure DAta (image, attributes, connection table and
          map table if it contains data).
NOS ----- NO Structure data.
ENTER STRUCTURE FORMAT (SIM), NOS:end

=> d L1
L1 HAS NO ANSWERS
L1              STR
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

Structure attributes must be viewed using STN Express query preparation.

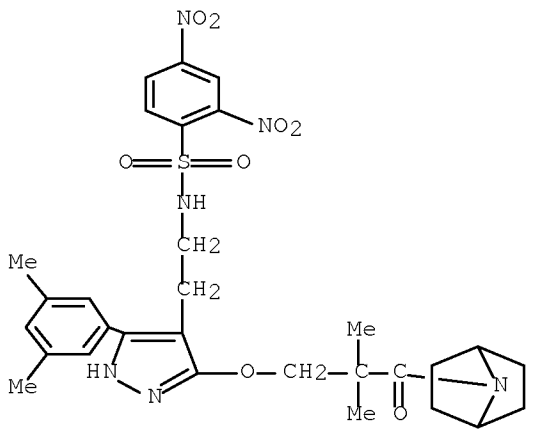

=> s L1 SSS full
FULL SEARCH INITIATED 13:16:44 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -    16968 TO ITERATE

100.0% PROCESSED    16968 ITERATIONS                           128 ANSWERS
SEARCH TIME: 00.00.01

L2          128 SEA SSS FUL L1

=> d scan

L2   128 ANSWERS   REGISTRY  COPYRIGHT 2008 ACS on STN
IN   7-Azabicyclo[2.2.1]heptane, 7-[3-[[5-(3,5-dimethylphenyl)-4-[2-[[(2,4-
     dinitrophenyl)sulfonyl]amino]ethyl]-1H-pyrazol-3-yl]oxy]-2,2-dimethyl-1-
     oxopropyl]- (9CI)
MF   C30 H36 N6 O8 S
```



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):o
'O' IS NOT VALID HERE

To display more answers, enter the number of answers you would like to see. To end the display, enter "NONE", "N", "0", or "END".
HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> file caplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	178.82	179.03

FILE 'CAPLUS' ENTERED AT 13:17:08 ON 15 APR 2008
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FILE COVERS 1907 - 15 Apr 2008 VOL 148 ISS 16
FILE LAST UPDATED: 14 Apr 2008 (20080414/ED)

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http://www.cas.org/infopolicy.html

=> d his

```
     (FILE 'HOME' ENTERED AT 13:15:38 ON 15 APR 2008)

     FILE 'REGISTRY' ENTERED AT 13:15:45 ON 15 APR 2008
L1             STRUCTURE UPLOADED
L2          128 S L1 SSS FULL

     FILE 'CAPLUS' ENTERED AT 13:17:08 ON 15 APR 2008

=> s L2 SSS full
L3            2 L2

=> s L1 SSS full
   REGISTRY INITIATED
Substance data SEARCH and crossover from CAS REGISTRY in progress...
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.




FULL SEARCH INITIATED 13:17:32 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -    16968 TO ITERATE

100.0% PROCESSED    16968 ITERATIONS                              128 ANSWERS
SEARCH TIME: 00.00.01

L4          128 SEA SSS FUL L1



L5            2 L4

=> file marpat
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                         ENTRY    SESSION
FULL ESTIMATED COST                                       0.48     358.35

FILE 'MARPAT' ENTERED AT 13:17:48 ON 15 APR 2008
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FILE CONTENT: 1961-PRESENT VOL 148 ISS 14 (20080411/ED)

SOME MARPAT RECORDS ARE DERIVED FROM INPI DATA FOR 1961-1987

MOST RECENT CITATIONS FOR PATENTS FROM MAJOR ISSUING AGENCIES
(COVERAGE TO THESE DATES IS NOT COMPLETE):

US  20080051413 28 FEB 2008
DE 102006039038 21 FEB 2008
EP      1889831 20 FEB 2008
JP   2008044933 28 FEB 2008
WO   2008028336 13 MAR 2008
GB      2440819 13 FEB 2008
FR      2904973 22 FEB 2008
RU      2317993 27 FEB 2008
CA      2593150 06 JAN 2008
```

Expanded G-group definition display now available.

Effective December 15th the iteration and answer limits in MARPAT have increased from 100,000 to 200,000 for both on-line and batch searches. For more information on MARPAT search limits, type HELP SLIMITS at an arrow prompt.

=> d his

(FILE 'HOME' ENTERED AT 13:15:38 ON 15 APR 2008)

FILE 'REGISTRY' ENTERED AT 13:15:45 ON 15 APR 2008
L1 STRUCTURE UPLOADED
L2 128 S L1 SSS FULL

FILE 'CAPLUS' ENTERED AT 13:17:08 ON 15 APR 2008
L3 2 S L2 SSS FULL
S L1

FILE 'REGISTRY' ENTERED AT 13:17:32 ON 15 APR 2008
L4 128 S L1 SSS FULL

FILE 'CAPLUS' ENTERED AT 13:17:33 ON 15 APR 2008
L5 2 S L4 SSS FULL

FILE 'MARPAT' ENTERED AT 13:17:48 ON 15 APR 2008

=> s L2 SSS full
FULL SEARCH INITIATED 13:18:03 FILE 'MARPAT'
FULL SCREEN SEARCH COMPLETED - 4276 TO ITERATE

100.0% PROCESSED 4276 ITERATIONS 5 ANSWERS
SEARCH TIME: 00.00.09

L6 5 SEA SSS FUL L1

=> file caplus
COST IN U.S. DOLLARS SINCE FILE ENTRY TOTAL SESSION
FULL ESTIMATED COST 64.96 423.31

FILE 'CAPLUS' ENTERED AT 13:18:26 ON 15 APR 2008
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FILE COVERS 1907 - 15 Apr 2008 VOL 148 ISS 16
FILE LAST UPDATED: 14 Apr 2008 (20080414/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply.
They are available for your review at:

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=> s L6
L7 5 L6

=> d ibib abs hitstr 1-
YOU HAVE REQUESTED DATA FROM 5 ANSWERS - CONTINUE? Y/(N):y

L7 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:1213081 CAPLUS Full-text
DOCUMENT NUMBER: 147:462329
TITLE: Methods for treating glaucoma and macular degeneration
INVENTOR(S): Rao, Vasantha P.; Casey, Patrick J.; Deng, Peifeng; Peterson, Yuri K.
PATENT ASSIGNEE(S): Duke University, USA
SOURCE: U.S. Pat. Appl. Publ., 26pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
US 20070249010	A1	20071025	US 2007-716724		20070312
PRIORITY APPLN. INFO.:			US 2006-781730P	P	20060314

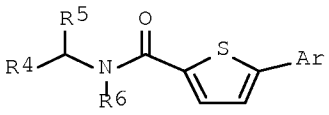
OTHER SOURCE(S): MARPAT 147:462329
AB Inhibition of geranylgeranylation of Rho GTPases and heterotrimeric G-proteins in the aqueous outflow pathway increases aqueous humor outflow, possibly through the tissue relaxation, and through altered cell adhesive interactions and actin cytoskeletal organization in cells of the outflow pathway. In addition, such inhibition is useful for treating age-related macular degeneration. The GGTase-I enzyme is a mol. target for lowering increased ocular pressure in glaucoma patients.

L7 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:493566 CAPLUS Full-text
DOCUMENT NUMBER: 141:38610
TITLE: Preparation of substituted thiophenes and related compounds as prenylation inhibitors
INVENTOR(S): Li, Francine Feirong; Rehder, Kenneth S.; Campbell, Michael Gordon; Viscardi, Celeste Patrice; Strachan, Jon-paul; Guo, Zhengming
PATENT ASSIGNEE(S): Ppd Discovery, Inc., USA
SOURCE: U.S. Pat. Appl. Publ., 117 pp., Cont.-in-part of U.S. Ser. No. 336,285.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 6
PATENT INFORMATION:

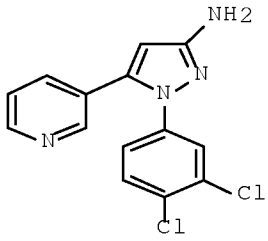
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20040116425	A1	20040617	US 2003-636327	20030806
US 7166619	B2	20070123		

US 6649638	B1	20031118	US 2003-336285	20030103
US 20070149549	A1	20070628	US 2007-618932	20070101
PRIORITY APPLN. INFO.:			US 2002-219628	B2 20020814
			US 2003-336285	A2 20030103
			US 2003-454554P	P 20030314
			US 2003-636327	A3 20030806

OTHER SOURCE(S): MARPAT 141:38610
GI



I



II

AB Title compds. I [Ar = heterocyclyl; R4 = absent, H, NH2, CONMe2, etc.; R5 = absent, i-Pr, benzyl, etc.; R6 = H, Me, Et, Pr, etc.] and related compds. are prepared For instance, 1-(3,4-dichlorophenyl)-5-(pyridin-3-yl)- 1H-pyrazole-3-carboxylic acid Me ester•HCl (preparation given) is saponified (THF/H2O, NaOH) and converted to the Boc-protected pyrazole-3-amine (i. DMF, t-BuOH, DPPA, Et3N; ii. t-BuOH, reflux, 4 h) and deprotected to II. Compds. of the invention have inhibitory activity for GTPase I [no data]. I inhibit protein prenylation and are useful for treating cancer, restenosis, psoriasis, etc.

L7 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER:	2004:182871 CAPLUS Full-text
DOCUMENT NUMBER:	140:235706
TITLE:	Preparation of substituted pyrazole derivatives as GnRH inhibitors
INVENTOR(S):	Bird, Thomas Geoffrey Colerick; Maudet, Mickael Louis Pierre
PATENT ASSIGNEE(S):	AstraZeneca AB, Swed.; AstraZeneca UK Limited
SOURCE:	PCT Int. Appl., 53 pp. CODEN: PIXXD2
DOCUMENT TYPE:	Patent
LANGUAGE:	English
FAMILY ACC. NUM. COUNT:	1
PATENT INFORMATION:	

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004018459	A1	20040304	WO 2003-GB3623	20030819

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

AU 2003255811	A1	20040311	AU 2003-255811	20030819
EP 1539743	A1	20050615	EP 2003-792484	20030819
EP 1539743	B1	20070530		

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK

JP 2006501231	T	20060112	JP 2004-530359	20030819
AT 363481	T	20070615	AT 2003-792484	20030819
ES 2286493	T3	20071201	ES 2003-792484	20030819
US 20050239858	A1	20051027	US 2005-525111	20050218
US 7253290	B2	20070807		

PRIORITY APPLN. INFO.: EP 2002-292075 A 20020821
WO 2003-GB3623 W 20030819

OTHER SOURCE(S): MARPAT 140:235706
GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Title compds. I [A = bond, alkylene; B = C(:X)R11; R1 = alkyl, etc.; R2 = mono- or bicyclic aromatic ring, etc.; R5 = 3-8 membered heterocyclic ring, etc.; R6-6a = H, alkyl, together equal CO; R7 = H, alkyl; X = N, CH; R8 = CN, H, OH, alkoxy, etc.; R11 = substituted amino, etc.] are prepared For instance, II (prepared in 4 steps from Me 3,5-dimethylbenzoate, butyrolactone and hydrazine) is reacted with 4-nitrophenyl chloroformate and 4-(pyrrolidin-3-yl)pyridine to give III. In an assay of GnRH-induced LH release, compds. of the invention are active at 1nM to 5μM.

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:182700 CAPLUS Full-text
DOCUMENT NUMBER: 140:235705
TITLE: Preparation of pyrazole derivatives as antagonists of gonadotropin releasing hormone (gnrh) for treating sex hormone related conditions
INVENTOR(S): Bird, Thomas Geoffrey Colerick; Herdemann, Matthias Ferdinand; Maudet, Mickael Louis Pierre
PATENT ASSIGNEE(S): AstraZeneca AB, Swed.; AstraZeneca UK Limited
SOURCE: PCT Int. Appl., 156 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

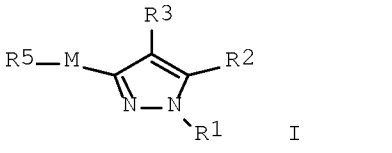
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004017961	A2	20040304	WO 2003-GB3633	20030819
WO 2004017961	A3	20040408		

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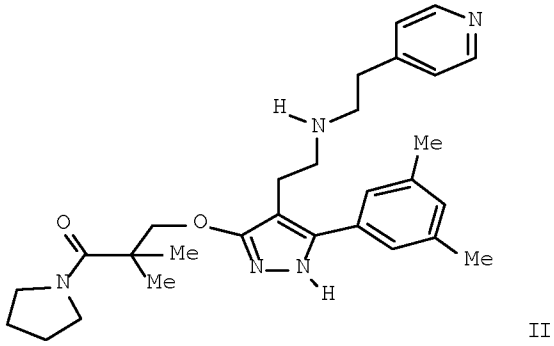
TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

AU 2003255820	A1	20040311	AU 2003-255820	20030819
EP 1531811	A2	20050525	EP 2003-792487	20030819
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JP 2006505528	T	20060216	JP 2004-530361	20030819
US 20060287379	A1	20061221	US 2005-524977	20051128
PRIORITY APPLN. INFO.:			EP 2002-292077 A	20020821
			WO 2003-GB3633 W	20030819

OTHER SOURCE(S): MARPAT 140:235705
GI



I



II

AB Title compds. I [R1 = H, (un)substituted-alkyl, aryl, or arylalkyl; R2 = (un)substituted mono or bicyclic aromatic ring; R3 = arylalkylaminoalkyl, heterocyclylalkylaminoalkyl, etc.; M = -(CH2)0-2O-, or -CONH-; R5 = H, halo, heterocyclylcarbonylalkyl, etc.] and their pharmaceutically acceptable salts are prepared and disclosed as gonadotropin releasing hormone antagonists. Thus, e.g., II, was prepared in a multistep synthesis from Me 3,5-dimethylbenzoate and butyrolactone. In test assays, I possessed activity at concns. from 1 nM to 5 μM.

L7 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

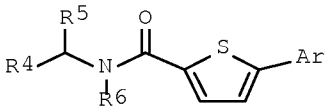
ACCESSION NUMBER:	2004:162671 CAPLUS Full-text
DOCUMENT NUMBER:	140:199323
TITLE:	Preparation of substituted thiophenes and related compounds as prenylation inhibitors
INVENTOR(S):	Li, Francine Feirong; Rehder, Kenneth S.; Campbell, Michael Gordon; Viscardi, Celeste Patrice; Strachan, Jon-Paul; Guo, Zhengming

PATENT ASSIGNEE(S): PPD Discovery, Inc., USA
SOURCE: PCT Int. Appl., 137 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 6
PATENT INFORMATION:

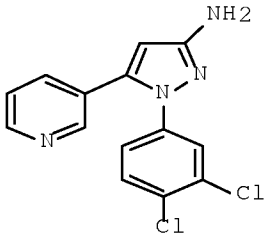
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004016592	A1	20040226	WO 2003-US24985	20030806
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 6649638	B1	20031118	US 2003-336285	20030103
AU 2003265395	A1	20040303	AU 2003-265395	20030806
EP 1534680	A1	20050601	EP 2003-788371	20030806
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				

PRIORITY APPLN. INFO.:		
US 2002-219628	A	20020814
US 2003-336285	A	20030103
US 2003-454554P	P	20030314
WO 2003-US24985	W	20030806

OTHER SOURCE(S): MARPAT 140:199323
GI



I



II

AB Title compds. I [Ar = heterocyclyl; R4 = absent, H, NH2, CONMe2, etc.; R5 = absent, i-Pr, Benzyl, etc.; R6 = H, Me, Et, Pr, etc.] and related compds. are prepared For instance, 1-(3,4-dichlorophenyl)-5-(pyridin-3-yl)- 1H-pyrazole-3-carboxylic acid Me ester•HCl (preparation given) is saponified (THF/H2O, NaOH) and converted to the Boc-protected pyrazole-3-amine (i. DMF, t-BuOH, DPPA, Et3N; ii. t-BuOH, reflux, 4 h) and deprotected to II. Compds. of the invention have inhibitory activity for GTPase I [no data]. I inhibit protein prenylation and are useful for treating cancer, restenosis, psoriasis, etc.

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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LOGOFF? (Y)/N/HOLD:y
STN INTERNATIONAL LOGOFF AT 13:18:55 ON 15 APR 2008
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